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Induction of heme oxygenase-1 contributes to survival of *Mycobacterium abscessus* in human macrophages-like THP-1 cellsMaher Y. Abdalla^{a,b,c,*}, Iman M. Ahmad^d, Barbara Switzer^{a,c}, Bradley E. Britigan^{a,b,c}^a Research Service, VA Medical Center—Omaha Nebraska Western Iowa, Omaha, NE 68105, USA^b Department of Pathology and Microbiology, University of Nebraska Medical Center College of Medicine, Omaha, NE 68198, USA^c Department of Internal Medicine, University of Nebraska Medical Center College of Medicine, Omaha, NE 68198, USA^d School of Allied Health Professions, University of Nebraska Medical Center College of Medicine, Omaha, NE 68198, USA

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ABSTRACT

Mycobacterium abscessus (M.abs) is a rapidly growing mycobacterial species that infects macrophages, and is an important pathogen in patients with cystic fibrosis. We studied the early stages of M.abs infection of macrophages, with emphasis on the role of heme-oxygenase-1 (HO-1) in this infection. THP-1 cells were activated using TPA into macrophage-like cells and infected with M.abs for different time points. M.abs infection robustly induced HO-1 expression in the THP-1 cells. Production of HO-1 was p38 MAPK-dependent, as p38 inhibitors suppressed HO-1 induction. Pretreatment with HO-1 inhibitors tin-protoporphyrin (SnPP) significantly inhibited M.abs growth inside macrophages. Furthermore, inhibiting HO-1 using HO-1 siRNA or the HO-1 upstream signaling molecule; Nrf2 using Nrf2 siRNA resulted in similar inhibition of M.abs. In contrast, inducing HO-1 did not increase M.abs intracellular growth above control. Products of HO-1 metabolism of heme are bilirubin, biliverdin, carbon monoxide (CO) and iron. The addition of either bilirubin or biliverdin, but not CO, completely restored the SnPP inhibitory effect and partially that with HO-1 siRNA. To understand the mechanisms, we used Syto-62 labeled M.abs to infect macrophages. Interestingly, HO-1 inhibition promoted M.abs-containing phagosome fusion with lysosomes, which should enhance M.abs killing. M.abs infection enhanced THP-1 ROS production as demonstrated by increased DHE, DCF fluorescence, and EPR signal. HO-1 inhibition further increased ROS production in infected macrophages. Our results indicate that HO-1 induction is important for M.abs growth during the early stages of infection, and that the HO-1 products bilirubin and biliverdin, perhaps through modulation of intracellular ROS levels, may be involved.

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Introduction

Mycobacterium abscessus (M.abs) is a rapidly growing non-tuberculous mycobacterial (NTM) species that infects macrophages of the lungs and skin and causes a variety of clinical syndromes in humans [1,2]. It has recently emerged as an important pathogen in patients with cystic fibrosis (CF), causing severe lung disease [3] and multiple complications that prevent lung transplantation [4]. Moreover, despite conventional cross-infection prevention procedures, frequent transmission of multidrug resistant NTM between patients with CF still exists [5].

Heme oxygenase-1 (HO-1) – also known as heat-shock protein 32 – is the rate-controlling enzyme of cellular heme catabolism. This microsomal enzyme acts on heme moieties to produce

equimolar amounts of carbon monoxide, iron (Fe), and biliverdin that is in turn converted to bilirubin by biliverdin reductase [6,7]. The Fe is then stored in ferritin, limiting its ability to participate as a catalyst through Fenton chemistry for production of cytotoxic free radicals [8]. Both biliverdin and bilirubin are thought to play an antioxidant role [9]. It was shown that HO-1 is induced by a variety of stimuli, such as ROS, viral infection and bacterial endotoxins, and appears to be protective in a variety of inflammatory disease states [10–12] due to its ability to inhibit inflammation and oxidative stress [13]. Moreover, induction of HO-1 suppresses apoptotic cell death through activation of MAPK and PI3K pathways with possible involvement of CO [14–17]. In THP-1 cells, HO-1 induction counteracted the effect of TNF-induced cell death via Nrf2 activation [18]. This is potentially of importance to mycobacterial infection as it appears that macrophage apoptosis contributes to host defense [19]. The role of CO in mycobacterial infection has been described previously. It was shown that *Mycobacterium tuberculosis* (M.tb) senses host-derived CO produced by HO-1 induction during macrophages infection [20], and CO

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activates the expression of dormancy (Dos) regulon [21], and other CO resistance genes such as *cor*, that protect bacteria from host-derived CO [22], and thus enhances mycobacterial survival *in vivo* [20–22]. We have also shown that acquiring Fe is critical to the metabolism and growth of mycobacteria and that limiting Fe availability could be a strategy for novel antimicrobial agents [23,24].

Little research has been conducted on *M. abs* pathogenesis. Like other pathogenic mycobacterial species, *M. abs* infects and multiplies within human macrophages, reflecting their ability to evade macrophage antimicrobial systems. Therefore, understanding the mechanism whereby *M. abs* successfully infect macrophages and finding approaches to reduce macrophage susceptibility to infection could lead to the development of novel antimicrobial agents to combat the growing clinical problem of multidrug resistant *M. abs*. The goal of this study is to assess the link between HO-1 and *M. abs* infection of human macrophages and understand the mechanisms by which HO-1 controls *M. abs* proliferation inside the infected macrophages.

Materials and methods

Materials

The p38 inhibitors; SB202190 hydrochloride was obtained from (Santa Cruz Biotechnology, Inc., Dallas, TX) and SB203580 from Cell Signaling Technology, Danvers, MA. The carbon monoxide (CO) releasing molecule (tricarbonylchloro(glycinato)ruthenium (II)) was purchased from Sigma, St. Louis, MO. Heme oxygenase-1 siRNA (h), Nrf2 siRNA (h), control siRNA, tin protoporphyrin IX dichloride, bilirubin, biliverdin hydrochloride, protoporphyrin IX cobalt chloride were all obtained from Santa Cruz Biotechnology.

Cell culture

THP-1 cells, a human monocyte-derived cell line, were obtained from the American Type Culture Collection (ATCC, Manassas, VA). The cells were grown in suspension in RPMI medium supplemented with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin (Thermo Scientific, Waltham, MA). The cells were cultured in 24-well tissue culture flask at a density of 10^6 cells/well in the presence of 15 ng/ml of 12-O-tetradecanoylphorbol-13-acetate (TPA; Cell Signaling Technology, Danvers, MA) for 24 h to differentiate the cells into a macrophage-like phenotype that adhere to the tissue culture plate. After infection with *M. abs* (MOI 5), the cells were placed in RPMI medium with 10% FBS and 30 μ g/ml gentamicin to inhibit extracellular replication of *M. abs*. To confirm that SnPP was not cytotoxic at the concentrations that were used, THP-1 cell viability was tested by flow cytometer in the presence of 7-aminoactinomycin D (7-AAD). No toxicity of SnPP was detected in infected or control cells at 24 h.

Cytotoxicity assay

Cytotoxicity was measured by the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) assay using CellTiter 96 Non-Radioactive Cell Proliferation Assay (Promega, Madison, WI).

Electron paramagnetic resonance spectroscopy

EPR spectroscopy was used to measure superoxide levels in *M. abs*-infected THP-1 cells. THP-1 cells were activated to macrophages and infected with *M. abs* for 1 h. Cells were then washed and incubated with antibiotic-free media for different time points. Treated uninfected cells were used as controls. Cells were

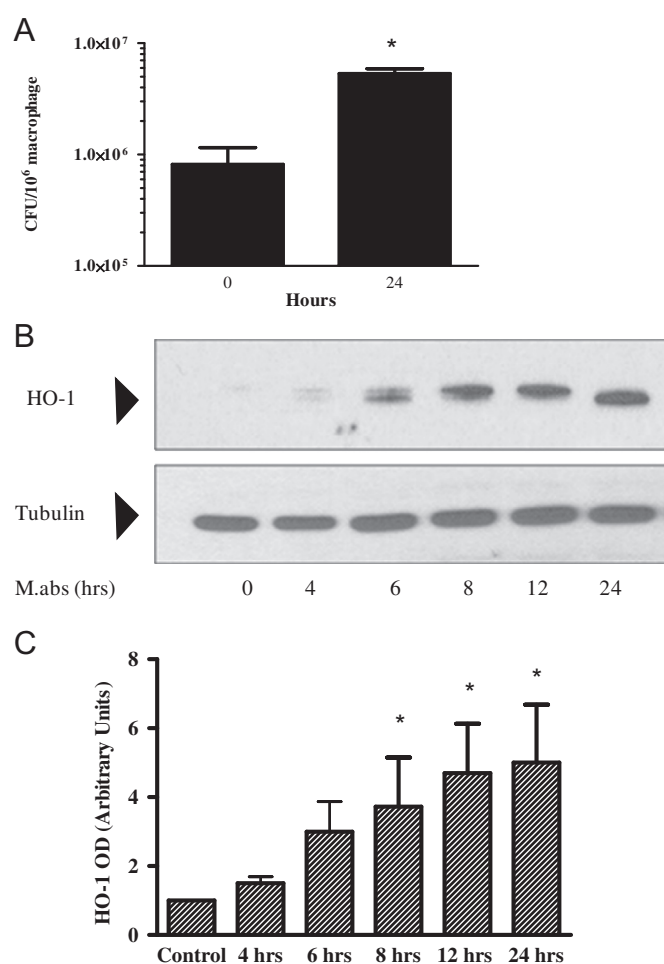


Fig. 1. *M. abs* increased HO-1 expression. TPA-differentiated THP-1 cells were infected with *M. abs* for 1 h, washed and incubated in media containing gentamicin until harvested at different time points. *M. abs* cfu were determined and cells were prepared for immunoblotting as described in [Materials and methods](#). (A) *M. abs* showed robust intracellular growth, as shown by the increase in CFU at days 0, and 1 post-infection. (B) Proteins were run and HO-1 visualized by immunoblotting. Results showed induction of HO-1 after infection. (C) Densitometric analysis of HO-1 bands in (B). *M. abs* infection resulted in a significant increase in expression of HO-1 ($n=4$, *indicates $p < 0.001$).

incubated (30 min at 37 °C) with CMH (200 μ M), a cell permeable, superoxide-sensitive spin probe in EPR buffer (pH 7.4). EPR buffer is a Krebs–HEPES buffer consisting of (in mM): 99 NaCl, 4.69 KCl, 2.5 CaCl₂, 1.2 MgSO₄, 25 NaHCO₃, 1.03 KH₂PO₄, 5.6 D-glucose, 20 HEPES and supplemented with the metal chelators DETC (5 μ M) and DF (25 μ M). After CMH incubation, macrophages were collected in a cell suspension, and 50 μ l of the cell suspension was placed into the Bruker Escan EPR spectrometer. The following EPR settings were used for all experiments: field sweep width, 60.0 G; microwave frequency, 9.75 GHz; microwave power, 21.90 mW; modulation amplitude, 2.37 G; conversion time, 10.24 ms; time constant, 40.96 ms. Levels of superoxide were normalized to the number of macrophages in each sample.

In situ ROS studies

In situ staining of superoxide (O₂^{•-}) and H₂O₂ levels were determined using the superoxide indicator dihydroethidium dihydroethidium (DHE) and the ROS indicator 5-(6)-chloromethyl-20,70-dichlorodihydrofluorescein diacetate (CM-H2DCFDA). *M. abs* bacteria were labeled with Syto-62 according to manufacturer's instruction (Invitrogen, Grand Island, NY). TPA-stimulated THP-1

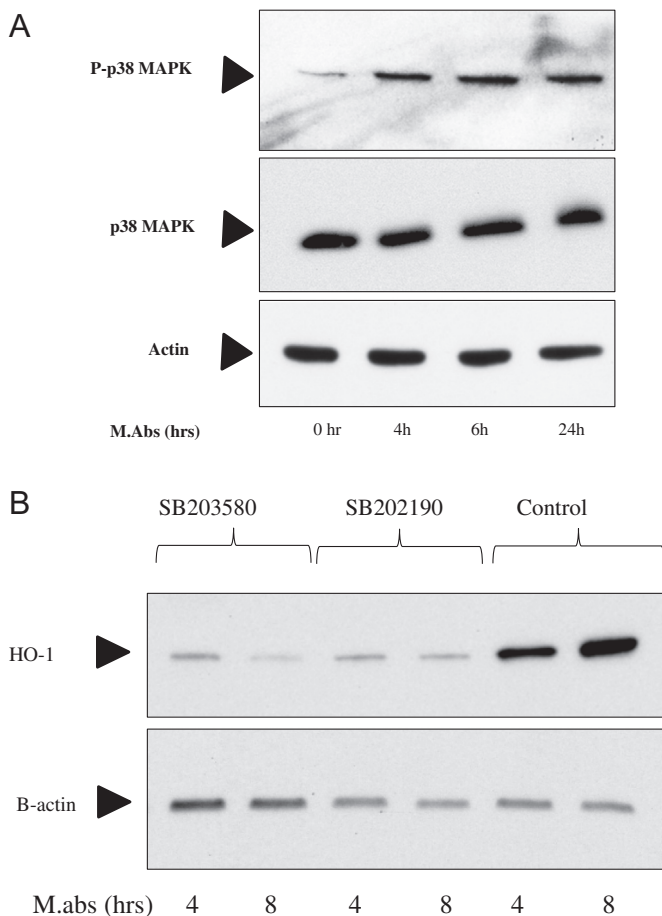


Fig. 2. p38 is involved in M.abs-mediated HO-1 induction signal transduction pathway. (A) TPA-differentiated THP-1 cells were infected with M.abs for 1 h, washed and incubated in media as shown in materials and methods. Cells were collected at different time points. Phosphorylated and total forms of p38MAPK were determined by immunoblotting. (B) Addition of the p38 inhibitors; SB203580 and SB202190 reduced HO-1 induction by M.abs.

cells were grown on a glass chamber slide and were infected with Syto-62-labeled M.abs for 1 h, and incubated with media for 4 h at CO₂ incubator. Thirty minutes before the infection was complete, DHE, and DCF were added to the assigned chambers. After infection was complete, the medium was removed, and chambers were washed, and mounted with Vectashield mounting medium with DAPI (Vector Laboratories, Burlingame, CA). Images were viewed using Zeiss 510 Meta Confocal Laser Scanning Microscope.

Western immunoblotting

Total protein lysates were prepared in RIPA buffer containing protease inhibitors (Thermo Scientific, Rockford, IL). Lysates were mixed with equal volume of 2 × Laemmli loading dye (Bio-Rad, Hercules, CA), boiled for 5 min at 95 °C, and loaded onto SDS-PAGE gels. After running, proteins were transferred to PVDF membranes, blocked with 5% milk in TBST, and probed with primary antibodies (p38 MAPK, Phospho-p38 MAPK, Cell Signaling Technology, Danvers, MA, and Anti-MnSOD, Anti-Catalase, Millipore, Billerica, MA) overnight at 4 °C with constant rocking. Membranes were then washed three times with TBST, incubated with secondary antibodies for 1 h at room temperature, washed three times with TBST, and protein were visualized using Pierce chemiluminescence reagents (Rockford, IL). Densitometry analyses were performed by NIH ImageJ.

Colony forming unit assay

Colony-forming units (CFUs) for M.abs present within THP-1 cells were determined as described previously [23]. In brief, cell culture supernatants were removed, and the plates were washed three times in sterile media. Chilled, chelexed, distilled water was added (0.3 ml/well) and plates were incubated on ice for 10 mins. After that, 1.2 ml lysis buffer (0.05% SDS, 5.0% BSA in Fe-free 7H9) was added to each well, and contents were scraped from plates, transferred to sterile tubes and centrifuged at 14,000g for 15 min at 4 °C. Bacteria were resuspended in 200 μl sterile Fe-free 7H9, and diluted for plating on tryptic soy agar plates (Remel, Lenexa, KS) for 3–5 days at 37 °C. The number of cfu on the plates was counted and the mean CFU was calculated.

Phagosome–lysosomes fusion studies

Macrophage-differentiated THP-1 cell were grown on a glass chamber slide and were infected with Syto-62 labeled M.abs for 1 h in antibiotics-free RPMI medium for 4 h. Two hours before the infection was complete, LysoTracker Red DND-99 (50 nM) was added to the cells to label the lysosomes (Invitrogen, Grand Island, NY). After infection was complete, the medium was removed and cells were washed and mounted with Vectashield mounting medium with DAPI (Vector Laboratories, Burlingame, CA). Co-localization and enumeration studies were performed on Zeiss 510 Meta Confocal Laser Scanning Microscope equipped with 4 lasers: a Blue Diode 405 nm; an Argon Laser 458/477/488,514 nm; a DPSS 561 nm and a HeNe 633 nm. Both, cells containing bacteria only and cells containing bacteria that co-localize with a lysosome were counted in controls and HO-1 inhibited cells. At least 50 cells were counted per condition in each experiment, and at least three independent experiments were performed. The results were expressed as % co-localization (number of cells that contain co-localized bacteria with lysosomes as percent of total cells counted).

SOD, catalase activity and GSH assays

Activity of antioxidant enzymes in macrophages was measured using the OxiSelect Catalase Activity Assay Kit (Cell Biolabs Inc, San Diego, CA) for catalase, GSSG/GSH Quantification kit (DOJINDO Inc, Rockville, MD) for total, oxidized and reduced GSH, and SOD Assay Kit-WST (DOJINDO Inc, Rockville, MD) to measure intracellular SOD activity according to the manufacturers' guidelines.

Analysis of data

Means and standard deviations were calculated from independent experiments. Statistical analysis was done using GraphPad Prism version 5 for Windows (GraphPad Software, San Diego, CA). Differences between three or more means were determined using one-way analysis of variance (ANOVA) with Bonferroni *post-hoc* tests. Error bars represent mean ± SEM. All statistical analyses were considered significant at $p < 0.05$. Each experiment was analyzed relative to its own control group(s).

Results

M.abs infection induces HO-1 expression in macrophages

The role of HO-1 in a variety of infections has been described before [12,25–28]. Therefore, we sought to study the level of HO-1 expression in M.abs infected and non-infected macrophages. To do so, we used the human monocytic cell line THP-1. Cells were differentiated into macrophages using TPA and then infected with M.

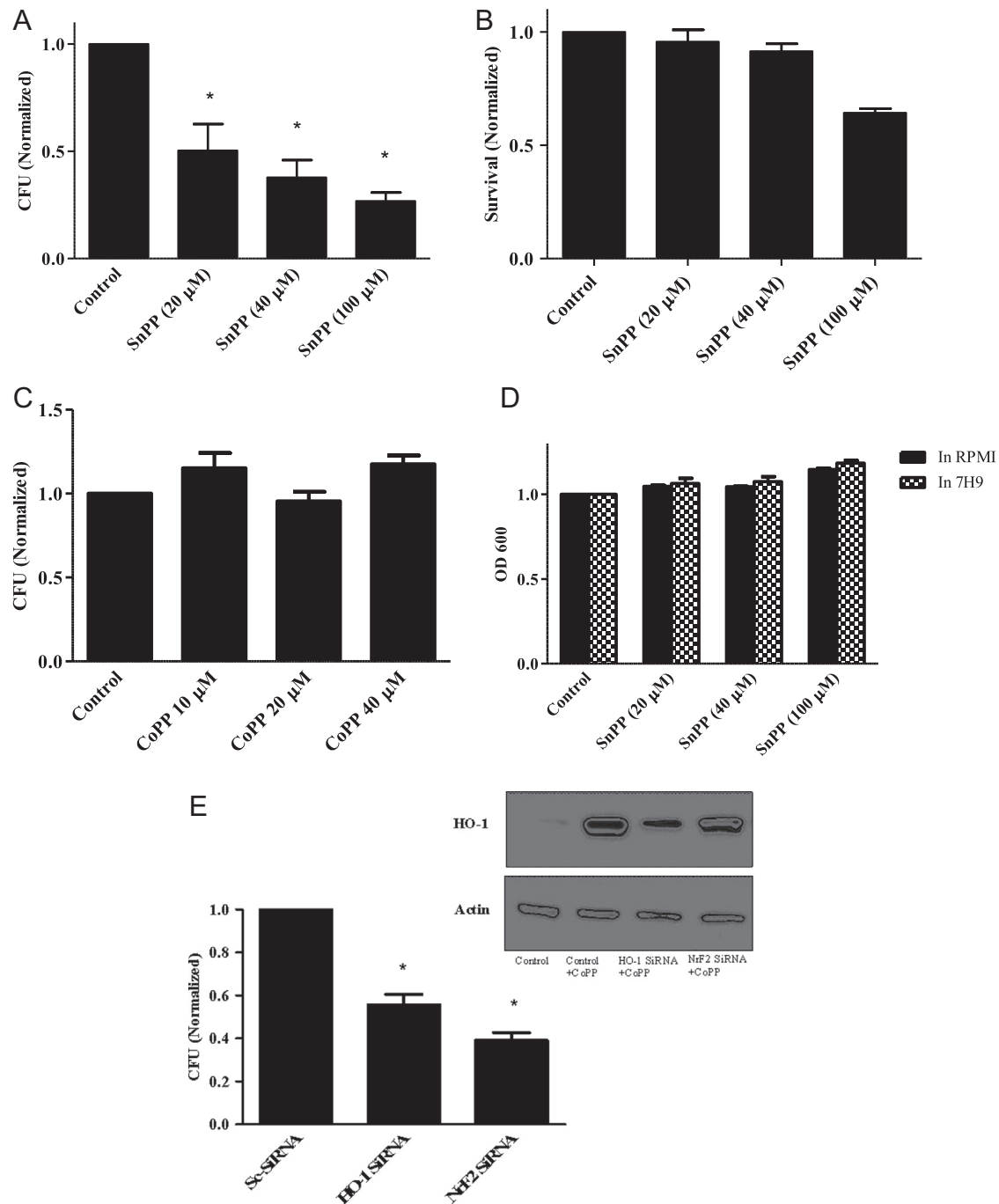


Fig. 3. HO-1 inhibition suppresses *M. abs* growth in macrophages. (A) TPA-differentiated THP-1 cells were cultured in the presence of the HO-1 inhibitor SnPP for 24 h, infected with *M. abs* for 1 h, and collected 24 h post-infection for CFU determination, as described in materials and methods. SnPP significantly inhibited *M. abs* growth as detected by CFU ($p < 0.05$, $n = 5$). Results obtained were normalized to vehicle-treated controls. (B) TPA-differentiated THP-1 cells were cultured in the presence of different concentrations of SnPP for 48 h and assayed for toxicity by MTT assay. No toxicity was seen using SnPP concentrations up to 40 μM for 48 h. Results were normalized to vehicle-treated controls. (C) No growth inhibition of *M. abs* was detected when bacteria were cultured in RPMI or 7H9 in the presence of different concentrations of SnPP. (D) TPA-differentiated THP-1 cells were cultured in the presence of different concentrations of CoPP for 24 h, and then infected with *M. abs* for 1 h. Cells were washed, and cultured in the presence of CoPP for 24 h. *M. abs* cfu were counted as described in materials and methods. There was no significant difference in *M. abs* growth in control versus CoPP treated THP-1 cells ($n = 3$, $p > 0.05$). CFU counts were normalized to vehicle-treated controls. (E) TPA-differentiated THP-1 cells were cultured in the presence of siRNA for 24 h and then infected with *M. abs* 1 hr. Cells were washed and cultured for 24 h before collected for CFU assay as described in materials and methods. Significant inhibition of *M. abs* growth was observed with treatment of THP-1 cells by HO-1 siRNA ($p < 0.05$, $n = 3$) or Nrf2 siRNA ($p < 0.05$, $n = 2$). CFU counts were normalized to Sc-siRNA-treated controls infected with *M. abs*. The inset shows decreased HO-1 protein level in the presence of HO-1 siRNA or Nrf2 siRNA. Cells were collected after 24 h of culture in the presence of HO-1 siRNA, Nrf2 siRNA, or CoPP (HO-1 inducer). Control cells treated with HO-1 siRNA shows no detectable protein by immunoblot (not shown). CoPP slightly induced HO-1 in the presence of siRNAs as determined by immunoblot ($p < 0.05$, $n = 3$).

abs. *M. abs* grew readily inside macrophages as shown by monitoring CFU at day 0 till day 1 (Fig. 1A). As shown in Fig. 1B, *M. abs* infection significantly increased HO-1 expression in a time-dependent manner, beginning 4 h post-infection ($p < 0.001$) (Fig. 1B, C).

M. abs induces HO-1 expression via p38 MAPK pathway

It has been shown that HO-1 induction is dependent on PI3K/p38MAPK signaling pathway in many cells, including THP-1 cells

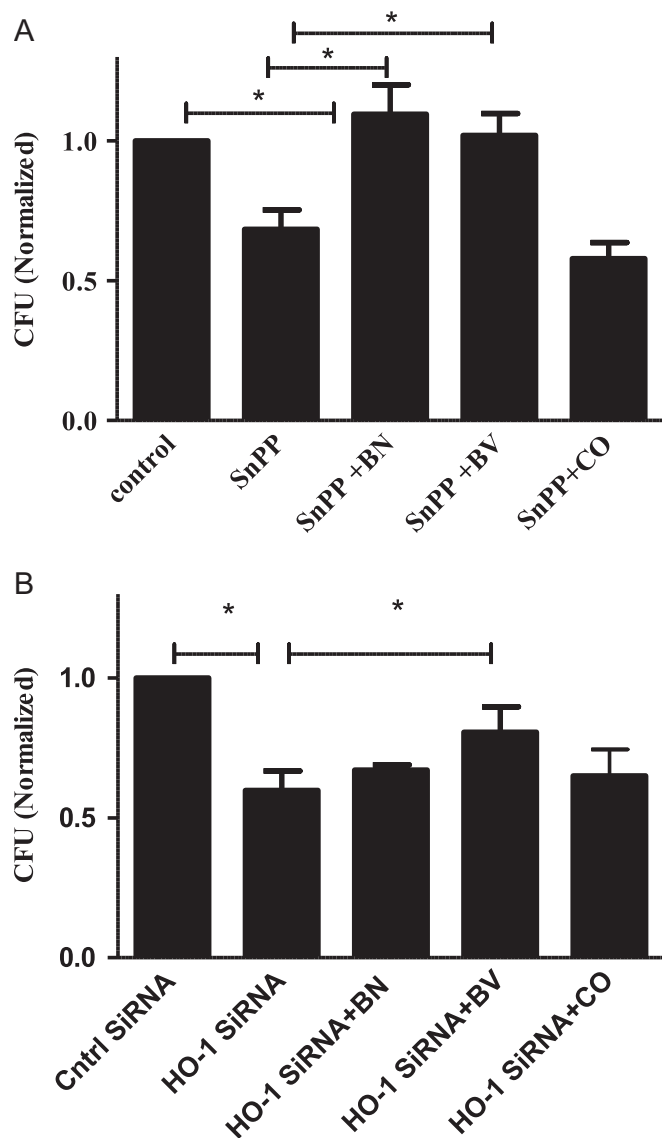


Fig. 4. HO-1 enzymatic products restore M.abs growth reduced by HO-1 inhibition. TPA-differentiated THP-1 cells were cultured in the presence of SnPP (30 μ M) or HO-1 siRNA for 24 h. Cells were then infected with M.abs and cultured in media with the addition of either bilirubin (10 μ M), biliverdin (10 μ M), or CO (10 μ M) for 24 h. Cells were collected and M.abs CFU assayed. CFU counts were normalized to vehicle-treated controls. (A) Both bilirubin and biliverdin significantly increased CFU in SnPP-treated cells ($n=7$, $p < 0.05$). (B) Biliverdin significantly increased CFU HO-1 siRNA-treated cells ($n=5$, $p < 0.05$).

[29,30]. When cells were collected at different times after M.abs infection, increased phosphorylated p38MAPK was shown starting at 4 h as shown in Fig. 2A. This is consistent with involvement of the p38MAPK signaling pathway in M.abs HO-1 induction of HO-1. Supporting this possibility, the administration of either of the p38 MAPK inhibitors SB203580 (10 μ M), or SB202190 (10 μ M) reduced M.abs-induced HO-1 protein expression (Fig. 2B).

HO-1 inhibition suppresses M.abs growth in macrophages

The induction of HO-1 could be very important for bacterial survival inside the macrophages. HO-1 induces Fe²⁺ storage as ferritin, thus limiting the generation of free radicals from free Fe²⁺ by Fenton reaction [8,31]. Moreover, both biliverdin and bilirubin play an antioxidant role that may contribute to the overall cytoprotective effect of HO-1 [9]. Iron is important for mycobacterial growth and virulence inside infected macrophages. Our laboratory

and others have shown that controlling iron availability could inhibit the growth of *Mycobacterium tuberculosis* and other bacteria growing within human macrophages [23,32–34].

Therefore, we examined the effect of altering HO-1 expression and/or HO-1 activity on M.abs growth inside infected macrophages. As shown in Fig. 3, inhibiting HO-1 activity with SnPP significantly ($p < 0.05$) decreased the number of M.abs CFU at 24 h, indicating suppression of bacterial growth in macrophages (Fig. 3A). The CFU determination was made at 24 h rather than 48 h because the infected THP-1 cells began to show evidence of detachment after 24 h, which could confound data interpretation.

It is important to mention that in these initial experiments, the SnPP was added to THP-1 cell cultures 24 h before infection, and added again after the infection until the time of cell collection. However, when SnPP was added at the time of infection, no inhibition of M.abs growth was observed (not shown). Furthermore, SnPP had no effect on M.abs phagocytosis as evidenced by a similar level of CFU on day 0 post-infection (data not shown). This suggests that the impact of HO-1 activity on M.abs growth is due to a process that occurs early after infection of the host cell.

To rule out any inhibitory effects of SnPP on M.abs or THP-1 cells, cells and bacteria were cultured in the presence of different concentrations of SnPP and monitored for toxicity. As shown in Fig. 3, no toxic effects were observed on either the THP-1 cells (B) or on the bacteria (C) when cultured with SnPP, as determined by the MTT test or OD600 absorbance, respectively. To further confirm our results and ensure that no toxic concentrations of SnPP were used, cells viability was assessed by flow cytometer in the presence of 7-AAD. No significant toxicity of SnPP, was seen in either control or M.abs-infected THP-1 cells at 24 h (not shown). Moreover, when similar experiments were performed in the presence of different concentrations of the HO-1 inducer CoPP, no significant increase in M.abs CFU were detected above untreated control, as shown in Fig. 3D ($p > 0.05$).

To further confirm our results, we inhibited HO-1 by HO-1 siRNA. Inhibiting HO-1 expression by HO-1 siRNA suppressed M.abs growth inside macrophage-differentiated THP-1 cells as demonstrated in Fig. 3E ($p < 0.05$). Moreover, inhibiting Nrf2, an upstream signaling factor for HO-1, resulted in a similar effect (Fig. 3E). Collectively, these results show HO-1 suppression at the time of infection inhibits M.abs growth in macrophages.

Impact of the HO-1 products bilirubin, biliverdin, and carbon monoxide on M.abs growth

The mechanism whereby M.abs HO-1 upregulation benefits M.abs intracellular survival was unclear. It has been shown that HO-1 products such as bilirubin, CO, and biliverdin reduce HCV replication *in vitro* [35,36]. Therefore, we sought to investigate the effect of HO-1 products on M.abs growth inside infected macrophages. In these set of experiments before infecting them with M.abs we inhibited THP-1 cell HO-1 by SnPP or HO-1 siRNA for 24 h to prevent any increase in HO-1 activity following M.abs infection. The cells were then infected with M.abs, and either bilirubin, biliverdin, or CO was added to the infected cells for another 24 h. Both bilirubin and biliverdin, but not CO, increased the number of CFU recovered from infected macrophages 24 h post-infection (Fig. 4). Bilirubin and biliverdin completely restored the CFU count to control levels with SnPP (Fig. 4A) and partially with HO-1 siRNA (Fig. 4B) treatment. The ability of bilirubin and biliverdin to restore CFU counts could be due to their antioxidant activity.

HO-1 inhibition increases phagosome–lysosome fusion

The phagocytosis of mycobacteria by macrophages is followed by phagosome–lysosome fusion [37,38]. The ability of

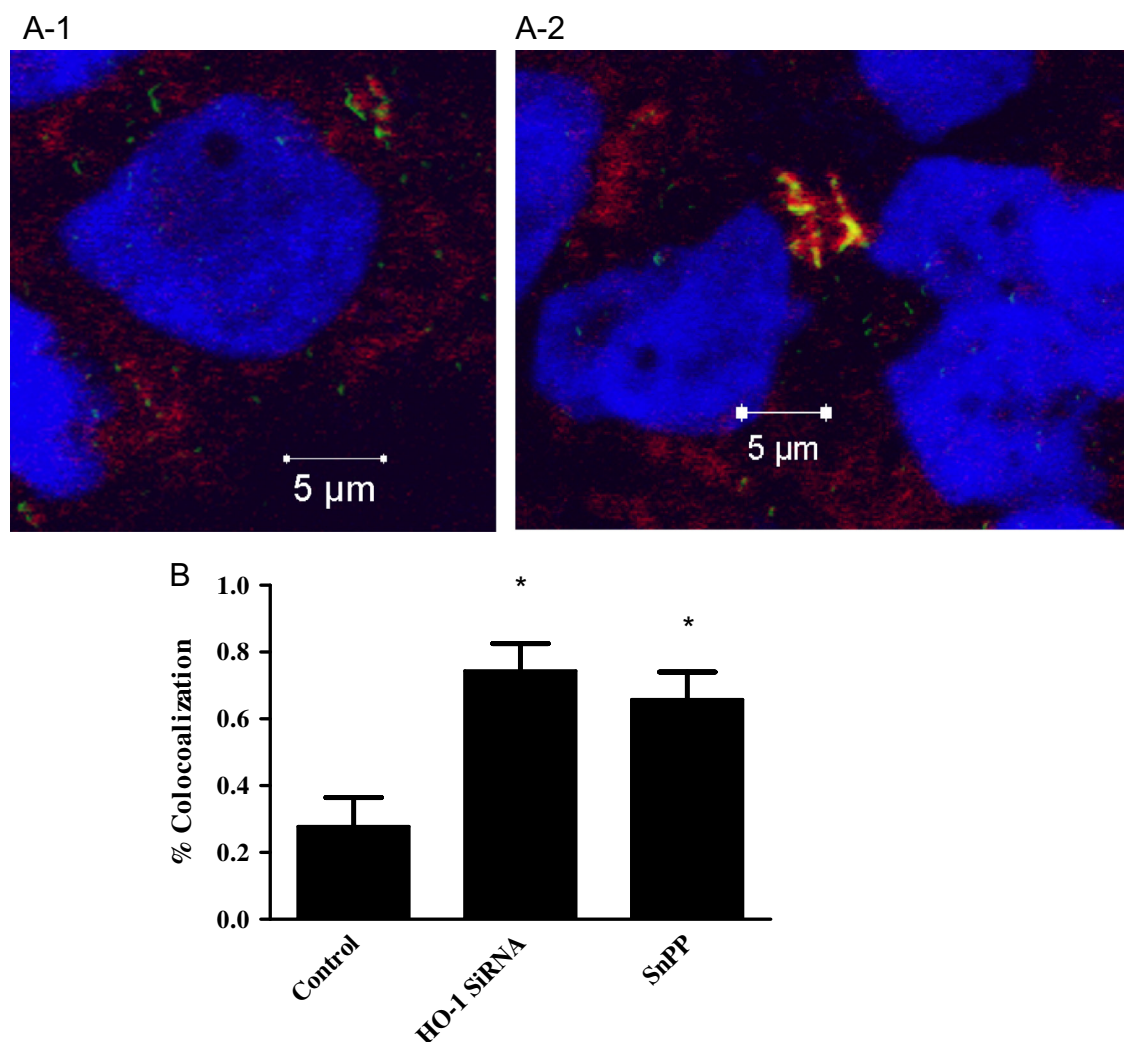


Fig. 5. HO-1 inhibition increases M.abs–lysosomal fusion in macrophages. TPA-differentiated THP-1 cells were cultured in the presence of SnPP or HO-1 siRNA for 24 h. Macrophages were infected with Syto-62 labeled M.abs for 1 h, and incubated for 4 h. LysoTracker was added 2 h before preparation for confocal microscopy. (A-1) control cells infected with M.abs (shown in green), lysotracker is shown in red, and nucleus in blue. (A-2) M.abs infection in the presence of HO-1 siRNA. M.abs–lysosomal co-localization is shown in yellow shows ($n=5$). (B) Percent co-localization in SnPP and HO-1 siRNA treated macrophages as compared to control. HO-1 inhibition resulted in a significant increase in the percentage of M.abs showing co-localization with lysosomes ($n=6$, $p < 0.05$).

mycobacteria to inhibit phagosome maturation into a phagolysosome is an important survival mechanisms used by pathogenic mycobacteria [39,40]. Therefore, we investigated whether HO-1 induction was involved in the ability of M.abs to inhibit phagosome–lysosome fusion. TPA-differentiated THP-1 cells were cultured in the presence of SnPP and HO-1 siRNA for 24 h. Macrophages were then infected for 4 h with Syto-62-labeled M.abs. Two hours before the completion of infection, LysoTracker was added to the cells to label the cells lysosomes. The cells were then viewed under a confocal microscope and the amount of co-localization of bacteria with lysosomes was calculated (Fig. 5). At 4 h after infection there was more co-localization observed in the macrophages with HO-1 inhibition as compared to controls (Fig. 5). This result is consistent with the possibility that HO-1 activity contributes to intracellular M.abs survival by playing a role in limiting phagosome–lysosome fusion.

M.abs infection induces oxidative stress and increases ROS production

HO-1 is also known to serve an anti-inflammatory role by decreasing intracellular oxidative stress. To assess the oxidative status of THP-1 cells infected with M.abs, TPA-differentiated THP-1

cells were infected with labeled M.abs for different time points. As shown in Fig. 6, when compared to non-infected control macrophages, M.abs infection increased intracellular superoxide ($O_2^{\bullet-}$) and H_2O_2 production as determined by DHE and DCFH2, respectively (Fig. 6). To confirm the increased ROS production, M.abs infected macrophages were tested by electron paramagnetic resonance (EPR) using the $O_2^{\bullet-}$ sensitive spin probe; CMH. As shown in Fig. 8B, a significant increase in CMH radical was detected after 4 h of M.abs infection, indicating more $O_2^{\bullet-}$ production ($p < 0.05$) (Fig. 8B).

Since, M.abs increased ROS production, we sought to assess the levels of antioxidant enzymes and compounds in M.abs infected macrophages. We first measured the levels of MnSOD and catalase protein levels in M.abs infected macrophages and controls at different time points. As shown in Fig. 7A, there were no differences between the levels of MnSOD and catalase protein over time (A). Similarly, there were no significant differences in either SOD (Fig. 7B) or catalase activity (C) at the different time points post-infection with M.abs ($p > 0.05$).

However, the picture was not the same when we measured the levels of reduced and oxidized GSH in M.abs infected and non-infected THP-1 cells. As observed in Fig. 7D, a decrease in the ratio

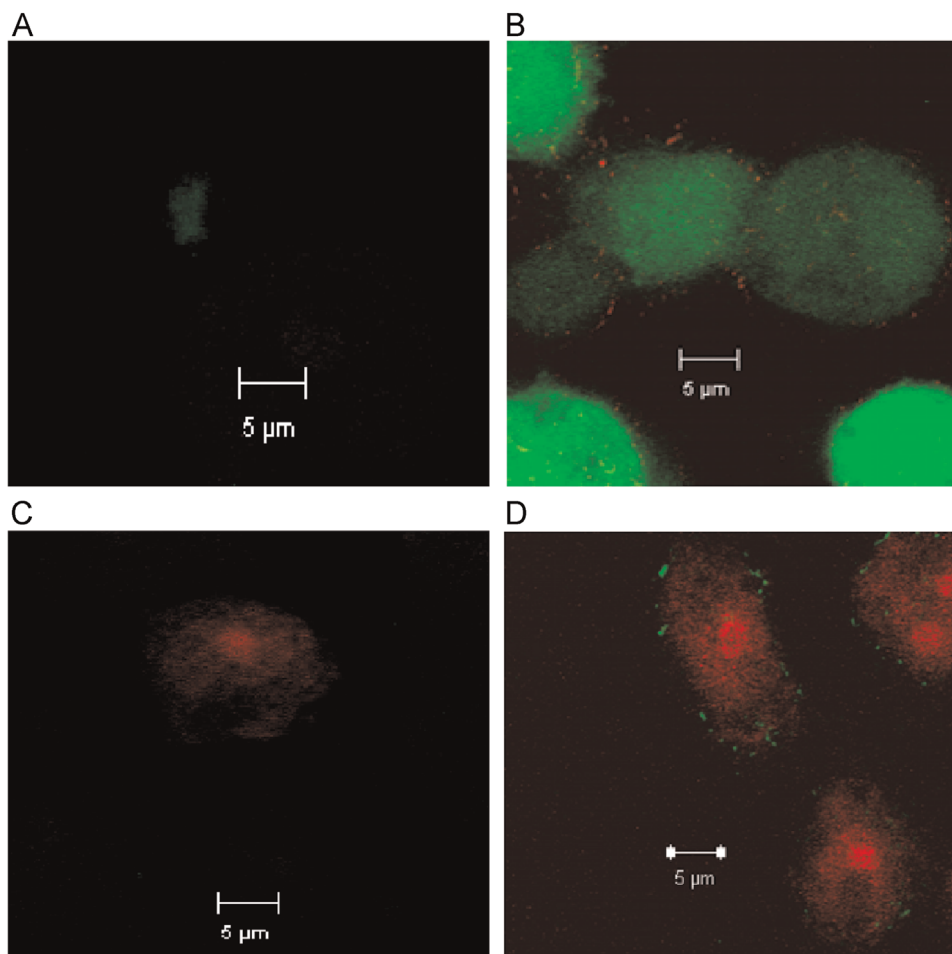


Fig. 6. M.abs infection increases THP-1 cell ROS production. TPA-differentiated THP-1 cells were infected with Syto-62 labeled M.abs for 4 h, and loaded with 10 μ M DCFH2 or DHE for the 30 min, washed and prepared for confocal microscopy study. Panels A and C are control cells with no infection showing minimal DCF or DHE fluorescence. Panel B shows increased DCF fluorescence and Panel D shows increased DHE fluorescence after 4 h of infection. Syto-62 labeled M.abs bacteria are shown inside the cells in red or green ($n=4$).

of reduced/oxidized glutathione (GSH/GSSG) ratio was detected after 4 and 6 h of infection compared to non-infecting controls, consistent with increased GSH turnover in M.abs infected cells as a result of increased oxidative stress (D).

HO-1 inhibition using SnPP increased ROS production as demonstrated by increased DHE and DCF fluorescence (Fig. 8A), and EPR measurements (B). Combining SnPP inhibition with M.abs infection led to more ROS production (Fig. 8B). This suggests that induction of HO-1 following M.abs infection of macrophages may serve to limit to levels of intracellular ROS.

Discussion

Mycobacterium abscessus is a rapidly growing mycobacterial species that can cause severe lung disease. Lung infections caused by M.abs are an emerging problem in the U.S. and many parts of the world, particularly in CF patients. Many M.abs strains are multi-drug resistant, and responsible for difficulties in patients undergoing lung transplantation [41]. M.abs infects and multiplies within human macrophages, indicating an ability of the bacteria to evade macrophage antimicrobial systems. In the current work, we sought to examine the early stages of M.abs infection inside macrophages, with a particular emphasis on the role of HO-1 in this process. We also defined the role of ROS and anti-oxidants inside macrophages infected with M.abs.

Our results show that M.abs infection increased $O_2^{\bullet-}$ and H_2O_2

production as indicated by increased fluorescence of DHE and DCF, respectively. Our EPR results confirmed the increase $O_2^{\bullet-}$ during the first hours of infection. Although protein levels and activities of SOD and catalase in the macrophages were not increased upon M.abs infection, reduced GSH was significantly decreased, indicating increased GSH turnover and oxidative stress. In contrast to SOD and catalase, HO-1 was induced during mycobacterial infection. This was not surprising as it was previously reported that HO-1 is induced in both infected macrophages and mice during *M. tuberculosis* infection [20,21]. HO-1 is highly inducible in response to a variety of stressful stimuli such as physical stress, reactive oxygen species (ROS), UV irradiation, endotoxins, and hypoxia and provide protection to cells and tissues against oxidative and inflammatory injury [10]. Induction of HO-1 is cytoprotective to host cells. Both biliverdin and bilirubin have antioxidant properties and can protect cells from a variety of ROS insults [9]. In addition, HO-1 prevents free Fe^{2+} accumulation within cell through sequestering it by ferritin and thus, limiting the generation of free radicals via the Fenton reaction [8,31]. CO produced by HO-1 has been reported to suppress apoptosis and modulate levels of pro-inflammatory cytokines, NO, and prostaglandins [42–44].

Since HO-1 is directly linked to macrophage Fe metabolism, HO-1 induction could contribute to pathogenesis by providing a new source of Fe for bacterial growth and replication. Alternatively, biliverdin/bilirubin production could improve the environment for M.abs growth. Finally, inhibition of macrophage apoptosis resulting from HO-1 induction could enhance M.abs

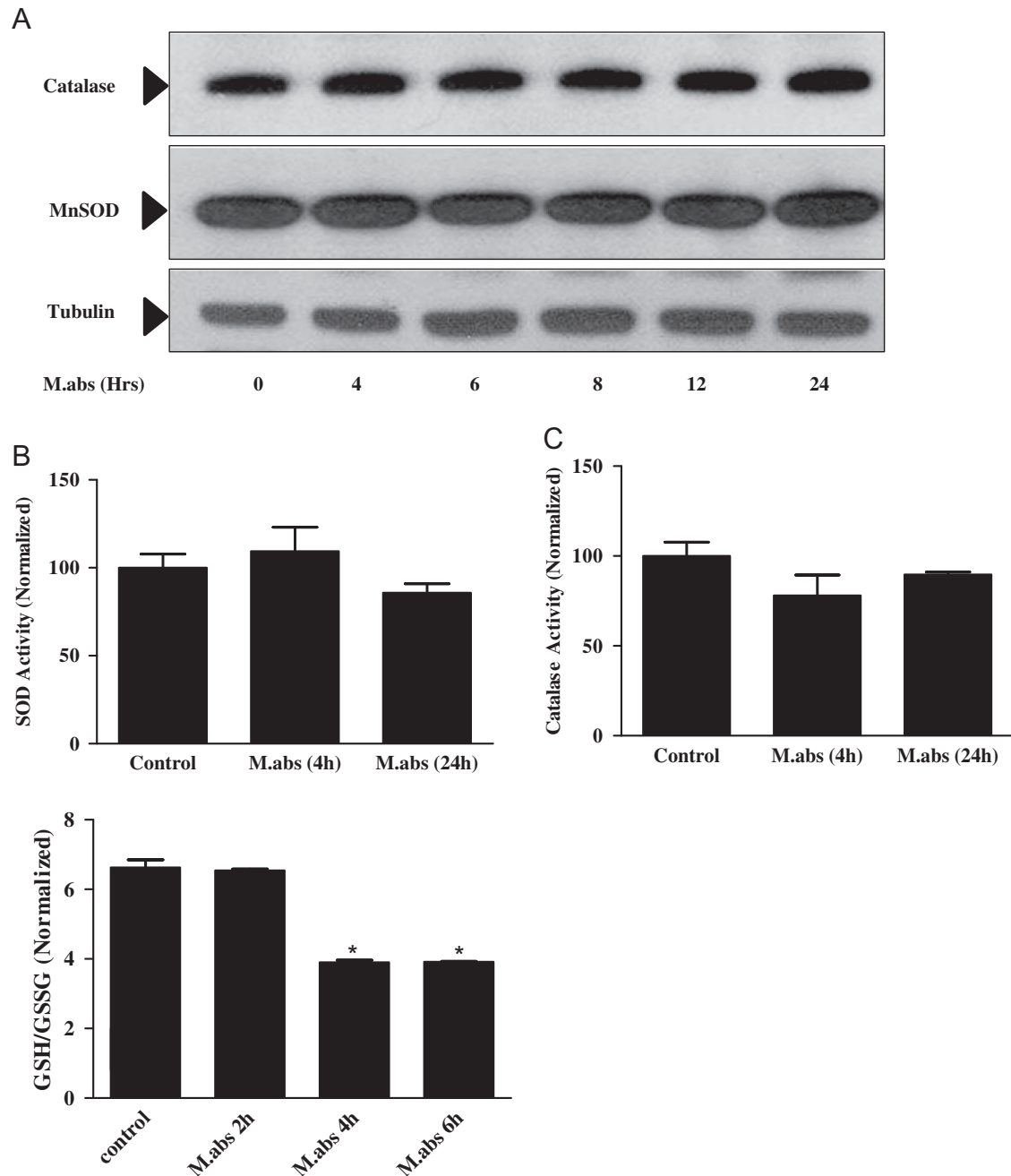


Fig. 7. M.abs infection increases THP-1 cell GSH turnover, but not MnSOD or catalase protein expression or activity. TPA-differentiated THP-1 cells were infected with M.abs for 1 h, washed and cultured in media until collected at different time points. Immunoblotting and enzyme activity were determined as described in materials and methods. (A) No difference in catalase or MnSOD protein (Panel A), SOD activity (Panel B, $n=3$, $p > 0.05$) or catalase activity (Panel C, $n=3$, $p > 0.05$) was observed between M.abs-infected and non-infected THP-1 cells at different points. (Panel D) Cells were then collected and studied for reduced glutathione (GSH), and oxidized glutathione (GSSG) levels as described in materials and methods. M.abs significantly decreased the ratio of GSH/GSSG, which reflects glutathione turnover ($n=3$, $p < 0.05$). Data counts in B–D were normalized to uninfected controls.

survival in macrophages. In fact, it was found that HO-1 favors pathogen dormancy in *M. tuberculosis* infections [21], and promotes the establishment of the malaria liver stage of infection [45]. Therefore, we assessed the importance of inhibiting HO-1 on mycobacterial growth inside infected macrophages.

We demonstrated that M.abs infection robustly induces HO-1 expression in the human macrophage-like THP-1 cells. Production of HO-1 was p38 MAPK-dependent, and p38 inhibitors suppressed HO-1 induction. We also demonstrated that pretreatment with HO-1 chemical inhibitors SnPP significantly inhibited M.abs growth inside macrophages, indicating the need of HO-1 for M.abs inside infected macrophages. It is worth mentioning that SnPP was

added 24 h before infection to inhibit HO-1, and adding SnPP at the time of infection did not show the same inhibitory effect. Furthermore, we showed that inhibiting HO-1 using HO-1 siRNA or the HO-1 upstream signaling molecule; Nrf2 using Nrf2 siRNA resulted in similar inhibitory effects. In contrast, inducing HO-1 using CoPP did not increase M.abs intracellular growth.

In seeming contrast to our findings, it has been shown that HO-1-deficient (*Hmox1*^{-/-}) mice were more susceptible to mycobacterial infection, and failed to mount a protective granulomatous response [46,47]. However, it is possible that the HO-1 effect in these studies extends beyond the macrophages and/or the lung alone since in murine models mycobacterial infection leads to

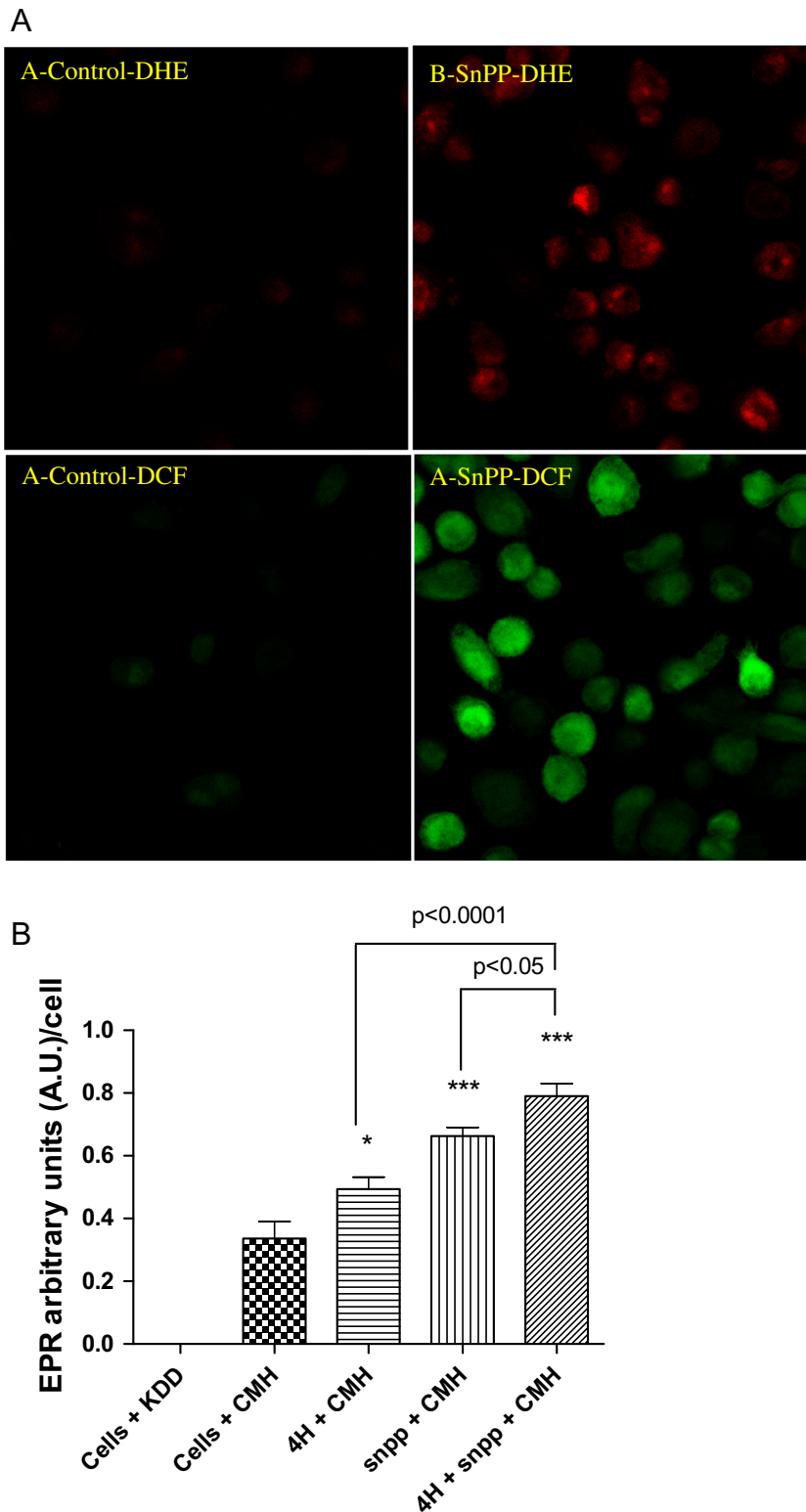


Fig. 8. HO-1 inhibition increase ROS production. TPA-differentiated THP-1 cells were infected with M.abs for 1 h, washed and cultured with media for 4 h. Cells were then loaded with DCFH2 or DHE for the 30 min, washed and prepared for confocal microscopy study. (A) HO-1 inhibition using SnPP increased ROS production as shown by DHE (A and B) and DCF fluorescence (C and D). (B) TPA-differentiated THP-1 cells were infected with M.abs for 4 h. Cells were then collected and studied for $O_2^{\bullet-}$ production by EPR as described in materials and methods. CMH is the spin probe 1-hydroxy-3-methoxycarbonyl-2,2,5,5-tetramethylpyrrolidine hydrochloride, KDD is the EPR buffer. A significant increase in $O_2^{\bullet-}$ production was observed at 4 h post-M.abs infection ($p < 0.05$), and when infection is combined with HO-1 inhibition ($p < 0.0001$). Inhibiting HO-1 also increased ROS production ($p < 0.0001$) ($n=5$).

systemic disease. Complete inhibition of HO-1 in whole body tissues might affect other functional roles of HO-1 in mycobacterial resistance.

Our data show that the presence of bilirubin and/or biliverdin reversed the negative impact of HO-1 inhibition on M.abs growth in THP-1 cells. This suggests that these HO-1 products may be

critical to the growth enhancing effect of HO-1 for M.abs in macrophages. The role of bilirubin and biliverdin as antioxidants extend beyond simply scavenging free radicals [48] to interfering with ROS production. It has been shown that decreased heme content due to HO-1 activation limits heme availability for the assembly of the functional NADPH oxidase subunits [49,50], and administration of the HO-1 inhibitor SnPP reversed this effect and increased NADPH [50]. The mechanism of inhibition could be partly mediated by bilirubin [50]. This is consistent with our results, where inhibiting HO-1 with SnPP increased ROS production as indicated by EPR and increased DHE and DCF fluorescence. The ability of bilirubin and biliverdin, but not CO, to reverse the effect of SnPP and increase M.abs growth inside macrophages supports a role for HO-1 activity and/or induction for M.abs survival inside macrophages. Further studies are still needed to define the exact mechanisms involved in enhancing M.abs intracellular growth.

The role of ROS in macrophage killing of intracellular mycobacteria is still debated. The enhanced bactericidal effect of increased ROS production in infected macrophages has been previously shown [51–53]. It has been suggested that *M. tuberculosis* may be more sensitive to ROS generation than other bacteria [52,53], and that minimizing the macrophage oxidative burst is one mechanism of mycobacterial intracellular survival [51]. On the other hand, it was reported that an oxidative environment could promote M.abs growth, and adding antioxidants and reducing intracellular oxidative stress could decrease intracellular growth of mycobacteria [54,55]. Combining our results with others, it could be concluded that both ROS and antioxidants are important for controlled, balanced intracellular environment that promotes mycobacterial killing and provides optimal conditions for functional macrophages against mycobacteria and other pathogens.

The effect of the HO-1 products could involve interaction with bacterial enzymatic activity and cellular functions. Iron inhibits the nonstructural 5B (NS5B) RNA-dependent RNA polymerase of HCV [56]. Moreover, biliverdin, and bilirubin were shown to inhibit HCV NS3/4A protease in liver cells and in cell-free assays [57,58].

A component of the intracellular survival mechanism of mycobacteria in human macrophages is the inhibition of macrophage phagosome–lysosome fusion [59,60]. Our results show that inhibiting HO-1 increased mycobacteria–lysosome co-localization and thus phagosome–lysosome fusion in M.abs infected macrophages. Increased phagosome–lysosome fusion in macrophages is important mechanism for mycobacterial killing. Mycobacteria have evolved multiple ways to evade or escape phagosome–lysosome fusion in macrophages. Tools for inhibiting phagosome–lysosome fusion ranged between producing ammonia in abundance [59], alkalizing the intralysosomal compartment [61], altered calmodulin-dependent signal transduction, and abundant iron availability [62]. In the latter, it was shown using dominant negative Rab5, that fusion of phagosomes with early endosomes and an adequate iron supply are important for mycobacteria to inhibit the phagosome maturation process [62]. This is consistent with our results that inhibiting HO-1 enhanced M.abs lysosomal co-localization, i.e. phagosome–lysosome fusion. This could be partially attributed to the reduced Fe availability that might result from lower HO-1 activity.

In summary, our results suggest that inhibiting HO-1 limits M.abs growth inside macrophages at the early stages of infection, and that reducing HO-1 products may enhance the ability of the macrophage to control M.abs infection. Further investigations are still required to define the detailed mechanisms involved and the potential for this to be translated to new forms of therapy against this emerging infection.

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